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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(use as many sheets as necessary)

Sheet 1 of 3

Complete if Known	
Application Number	09/936,957
Filing Date	September 17, 2001
First Named Inventor	Meikle
Group Art Unit	Unassigned
Examiner Name	Unassigned
Attorney Docket Number	021385-014010US

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	U.S. Patent Document Number	Kind Code[2] (if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
DC	AA	5,501,957		Dennis et al.	03-26-1996	

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Foreign Patent Document Office[3]	Number[4]	Kind Code[5] (if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T[6]

Examiner Signature		Date Considered	3/17/08

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 2 of 3

Complete if Known	
Application Number	09/936,957
Filing Date	September 17, 2001
First Named Inventor	Meikle
Group Art Unit	Unassigned
Examiner Name	Unassigned
Attorney Docket Number	021385-014010US

OTHER PRIOR ART -- NON PATENT LITERATURE DOCUMENTS

Examiner Initials*	Cite No.[1]	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T[2]
	AB	AERTS et al., "The occurrence of two immunologically distinguishable β-glucocerebrosidases in human spleen," Eur. J. Biochem., 150:565-574 (1985).	
	AC	BAGHDIGUIAN et al., "Co-localization of suramin and serum albumin in lysosomes of suramin-treated human colon cancer cells," Cancer Letters, 101:179-184 (1996).	
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	AJ	KISHIMOTO et al., "Saposins: structure, function, distribution, and molecular genetics," J. Lipid Research, 33:1255-1267 (1992).	
	AK	MEIKLE et al., "Diagnosis of lysosomal storage disorders: evaluation of lysosome-associated membrane protein LAMP-1 as a diagnostic marker," Clinical Chemistry, 43(8):1325-1335 (1997).	
	AL	MICHELAKAKIS et al., "Characterization of glucocerebrosidase in Greek Gaucher disease patients: mutation analysis and biochemical studies," J. Inher. Metab. Dis., 18:609-615 (1995), with abstract	
	AM	PASCHKE et al., "Infantile type of sialic acid storage disease with sialuria," Clinical Genetics, 29:417-424 (1986).	
	AN	RENLUND, M., "Clinical and laboratory diagnosis of Salla disease in infancy and childhood," Journal of Pediatrics, 104(2):232-236 (1979).	
	AO	RENLUND et al.., "Increased Urinary Excretion of Free *N*-Acetylneuraminic Acid in Thirteen Patients with Salla Disease," European Journal of Biochemistry, 101:245-250 (1979).	
	AP	RENLUND et al., "Salla disease: A new lysosomal storage disorder with disturbed sialic acid metabolism," Neurology, 33:57-66 (1983).	

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Sheet 3 of 3

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	AQ	RENLUND et al, "Studies on the Defect Underlying the Lysosomal Storage of Sialic Acid in Salia Disease: Lysosomal Accumulation of Sialic Acid Formed From *N*-Acetyl-Mannosamine or Derived from Low Density Lipoprotein in Cultured Mutant Fibroblasts," Journal of Clinical Investigation, 77:568-574 (1986).	
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	AT	WAHEED et al., "Enhanced Breakdown of Arylsulfatase A in Multiple Sulfatase Deficiency," European Journal of Biochemistry, 123:317-321 (1982).	

Examiner Signature		Date Considered	3/17/08

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